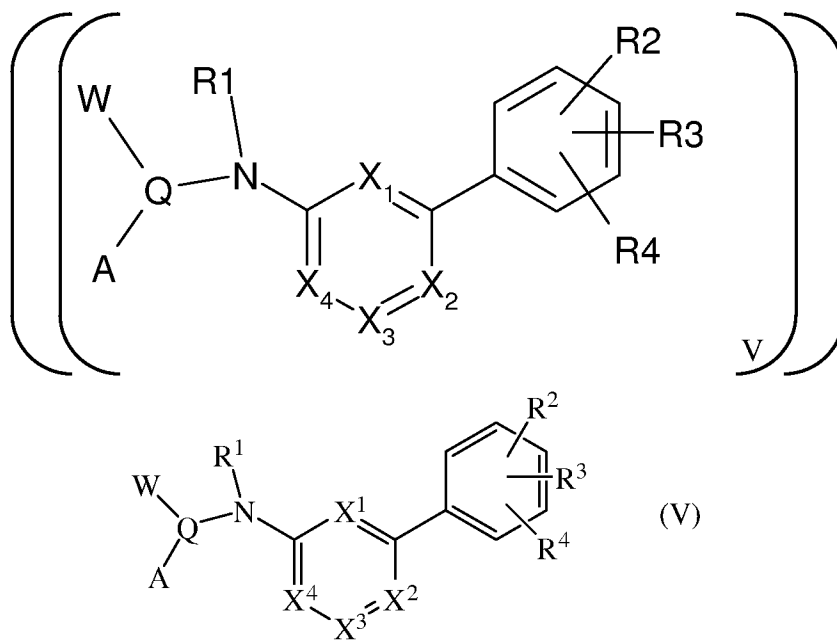


CLAIM AMENDMENTS

1-9. (canceled)

10. (currently amended): A compound of the ~~general~~ formula (V)

or pharmaceutically acceptable prodrugs, salts, hydrates, solvates, crystal forms, enantiomers, or diastereomers thereof, wherein X^1 and X^2 are N and X^3 and X^4 are C independently substituted with Y; [[:]]

~~X_1, X_2, X_3, X_4 are selected from the following:~~

- ~~(i) — X_1 and X_2 are N and X_3 and X_4 are C independently substituted with Y;~~
- ~~(ii) — X_1 and X_4 are N and X_2 and X_3 are C independently substituted with Y;~~
- ~~(iii) — X_2 and X_4 are N and X_1 and X_3 are C independently substituted with Y;~~
- ~~(iv) — X_1 is N and X_2, X_3 , and X_4 are C independently substituted with Y;~~
- ~~(v) — X_3 is N and X_1, X_2 , and X_4 are C independently substituted with Y;~~
- ~~(vi) — X_4 is N and X_1, X_2 , and X_3 are C independently substituted with Y;~~
- ~~(vii) — X_2 is N and X_1, X_3 , and X_4 are C independently substituted with Y; and~~
- ~~(viii) — X_1, X_2 and X_3 are N and X_4 is C substituted with Y;~~

[[R1]] R¹ is H, C₁₋₆ alkyl, ~~C₁₋₆ alkylNR⁵R⁶, C₁₋₆ alkylNR⁵COR⁶, C₁₋₆ alkylNR⁵SO₂R⁶, C₁₋₆ alkylCO₂R⁵, C₁₋₆ alkylCONR⁵R⁶, where R⁵ and R⁶ C₁₋₆ alkylNR⁵R⁶, C₁₋₆ alkylNR⁵COR⁶, C₁₋₆ alkylNR⁵SO₂R⁶, C₁₋₆ alkylCO₂R⁵, or C₁₋₆ alkylCONR⁵R⁶,~~

wherein R⁵ and R⁶ are each independently H, C₁₋₄ alkyl, aryl, hetaryl, C₁₋₄ alkylaryl, or C₁₋₄ alkylhetaryl or may be joined to form an optionally substituted 3-8 membered ring optionally containing ~~an atom selected from O, S, NR⁷~~ one of O, S or NR⁷;

~~R⁷ is selected from H, wherein R⁷ is H or C₁₋₄ alkyl;~~

[[R2]] R² is selected from OH, OC₁₋₆ alkyl, C₁₋₆ alkylOH, OC₂₋₆ alkylOH, ~~C₁₋₆ alkylNR⁸R⁹, OC₂₋₆ alkylNR⁸R⁹, C₁₋₆ alkylNR⁸COR⁹, OC₂₋₆ alkylNR⁸COR⁹, C₁₋₆ alkylNR⁸R⁹, OC₂₋₆ alkylNR⁸R⁹, C₁₋₆ alkylNR⁸COR⁹, OC₂₋₆ alkylNR⁸COR⁹, C₁₋₆ alkylhetaryl, OC₂₋₆ alkylhetaryl, CONR⁸R⁹, NR⁸COOR⁹, NR¹⁰CONR⁸R⁹, CONR⁸R⁹, NR⁸COR¹² OCONR⁸R⁹, NR⁸COOR⁹, NR¹⁰CONR⁸R⁹, CONR⁸R⁹, and NR⁸COR¹²;~~

~~R⁸, R⁹ wherein R⁸, R⁹ are each independently H, C₁₋₄ alkyl, C₁₋₄ alkylNR¹¹R¹³, C₁₋₄ alkylNR¹¹R¹³, hetaryl, or cyclohetalkyl, or may be joined to form an optionally substituted 3-8 membered ring optionally containing~~ an atom selected from O, S, NR¹⁴ ~~one of O, S or NR¹⁴;~~

[[R12]] wherein R¹² is C₂₋₄ alkyl, ~~C₁₋₄ alkylNR¹¹R¹³~~ C₁₋₄ alkylNR¹¹R¹³, hetaryl, or cyclohetalkyl;

~~R¹¹, R¹³ wherein R¹¹, and R¹³ are each independently H, or C₁₋₄ alkyl, or may be joined to form an optionally substituted 3-8 membered ring optionally containing~~ an atom selected from O, S, NR¹⁴ ~~one of O, S or NR¹⁴;~~

~~R¹⁴ is selected from H, wherein R¹⁴ is H or C₁₋₄ alkyl;~~

~~R¹⁰ is H, wherein R¹⁰ is H or C₁₋₄ alkyl;~~

~~R³ and R⁴~~ R³ and R⁴ are each independently H, halogen, C₁₋₄ alkyl, OH, OC₁₋₄ alkyl, CF₃, or OCF₃;

Q is a bond when W is absent, or C₁₋₄ alkyl when W is present;

W is selected from H, C₁₋₄ alkyl, and C₂₋₆ alkenyl; where C₁₋₄ alkyl or C₂₋₆ alkenyl may be optionally substituted with C₁₋₄ alkyl, OH, OC₁₋₄ alkyl, ~~NR¹⁵R¹⁶~~ or NR¹⁵R¹⁶;

~~R¹⁵, and R¹⁶~~ wherein R¹⁵, and R¹⁶ are each independently H, C₁₋₄ alkyl, C₁₋₄ alkyl cycloalkyl, C₁₋₄ alkyl cyclohetalkyl, aryl, or hetaryl, or may be joined to form an optionally

substituted 3-8 membered ring optionally containing ~~an atom selected from O, S, NR¹⁷~~ one of O, S or NR¹⁷;

[[R¹⁷]] wherein R¹⁷ is ~~selected from H, or C₁₋₄ alkyl~~;

A is aryl[[,]] or hetaryl optionally substituted with 0-3 substituents independently ~~chosen~~ selected from halogen, C₁₋₄ alkyl, CF₃, aryl, hetaryl, OCF₃, OC₁₋₄ alkyl, ~~OC₂₋₅ alkylNR¹⁸R¹⁹~~ OC₂₋₅ alkylNR¹⁸R¹⁹, Oaryl, Ohetaryl, ~~CO₂R¹⁸, CONR¹⁸R¹⁹, NR¹⁸R¹⁹, C₁₋₄ alkylNR¹⁸R¹⁹, NR²⁰C₁₋₄ alkylNR¹⁸R¹⁹, NR¹⁸COR¹⁹, NR²⁰CONR¹⁸R¹⁹, NR¹⁸SO₂R¹⁹~~ CO₂R¹⁸, CONR¹⁸R¹⁹, NR¹⁸R¹⁹, C₁₋₄ alkylNR¹⁸R¹⁹, NR²⁰C₁₋₄ alkylNR¹⁸R¹⁹, NR¹⁸COR¹⁹, NR²⁰CONR¹⁸R¹⁹, and NR¹⁸SO₂R¹⁹;

~~R¹⁸, R¹⁹~~ wherein R¹⁸ and R¹⁹ are each independently H, C₁₋₄ alkyl, C₁₋₄ alkyl cyclohetalkyl, aryl, hetaryl, C₁₋₄ alkyl aryl, or C₁₋₄ alkyl hetaryl, or may be joined to form an optionally substituted 3-8 membered ring optionally containing ~~an atom selected from O, S, NR²¹~~ one of O, S or NR²¹;

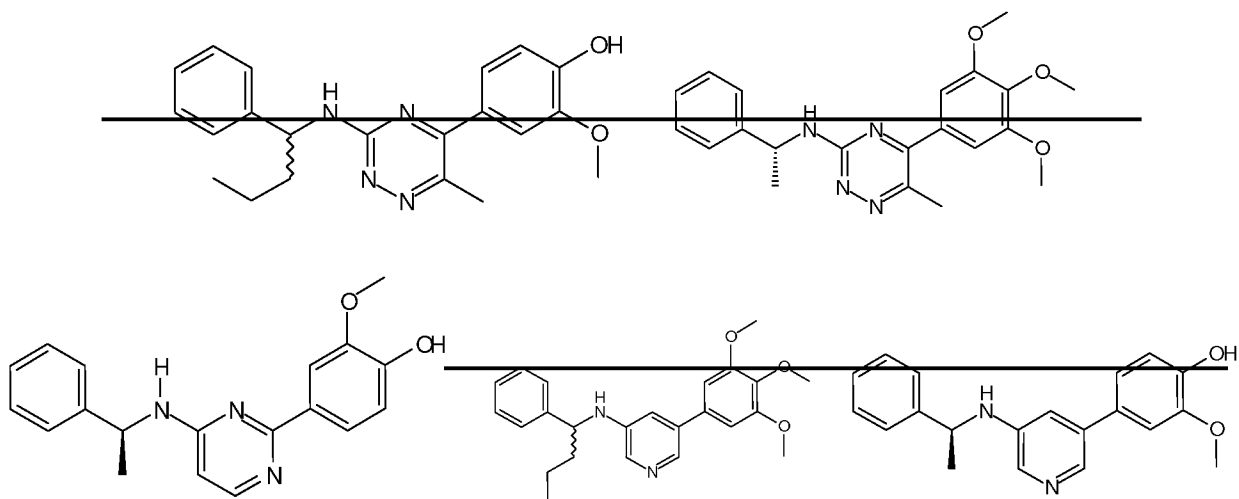
~~R²¹ is selected from H,~~ wherein R²¹ is H or C₁₋₄ alkyl;

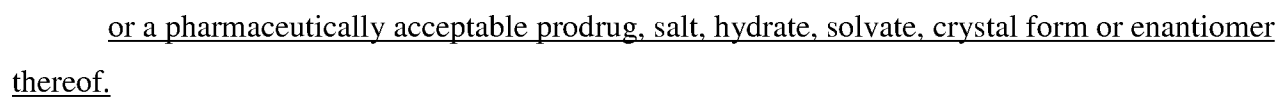
~~R²⁰ is selected from H,~~ wherein R²⁰ is H or C₁₋₄ alkyl;

Y is selected from H, C₁₋₄ alkyl, OH, ~~NR²²R²³~~ and NR²²R²³;

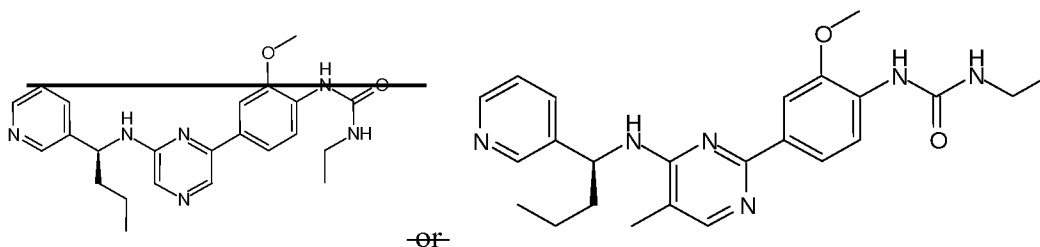
~~R²², R²³~~ wherein R²², R²³ are each independently H[[,]] or C₁₋₄ alkyl.

11. (currently amended): A compound according to claim 10 selected from the group consisting of:





12. (currently amended): A compound of the formula:



or a pharmaceutically acceptable prodrug, salt, hydrate, solvate, crystal form or enantiomer thereof.

13. (canceled)

14. (currently amended): A composition comprising a carrier and at least one compound according to ~~claim 1~~ claim 10.

15. (withdrawn; currently amended): A method ~~of treatment of~~ to treat a hyperproliferation-related disorder or disease state in a subject, said method comprising administering a therapeutically effective amount of at least one compound according to ~~claim 1~~ claim 10.

16. (withdrawn; currently amended): ~~A method of treatment according to~~ The method of claim 15, wherein the hyperproliferation-related disorder or disease state is treatable by the modulation of microtubule polymerisation.

17. (withdrawn; currently amended): ~~A method according to~~ The method of claim 15, wherein the hyperproliferation-related disorder or disease state is selected from the group consisting of ~~Cancer~~ cancer, infectious diseases, vascular restenosis or inflammatory diseases.

18. (withdrawn; currently amended): A method ~~of treatment of~~ to treat a protein-kinase related disorder or disease state in a subject, said method comprising administering a therapeutically effective amount of at least one compound according to ~~claim 1~~ claim 16.

19. (withdrawn; currently amended): ~~A method according to~~ The method of claim 18, wherein the protein-kinase related disorder or disease state is selected from the group consisting of ~~Atopy, Cell Mediated Hypersensitivity, Rheumatic Diseases, Other~~ atopy, cell mediated hypersensitivity, rheumatic diseases, other autoimmune diseases and ~~Viral Diseases~~ viral diseases.

20. (currently amended): A method ~~of treatment of~~ to treat diseases and conditions associated with inflammation and infection in a subject, said method comprising administering a therapeutically effective amount of at least one compound according to ~~claim 1~~ claim 10.

21. (new): A composition comprising a carrier and at least one compound according to claim 11.

22. (new): A composition comprising a carrier and at least one compound according to claim 12.

23. (new): The compound of claim 10, wherein R^2 is selected from C_{1-6} alkylOH, OC_{2-6} alkylOH, C_{1-6} alkylNR⁸R⁹, OC_{2-6} alkylNR⁸R⁹, C_{1-6} alkylNR⁸COR⁹, OC_{2-6} alkylNR⁸COR⁹, C_{1-6} alkylhetaryl, OC_{2-6} alkylhetaryl, $OCONR^8R^9$, NR^8COOR^9 , $NR^{10}CONR^8R^9$, $CONR^8R^9$, and NR^8COR^{12} , wherein R^8 , R^9 and R^{12} are as defined in claim 10.

24. (new): The compound of claim 23, wherein:
 R^1 is H, C_{1-6} alkyl, C_{1-6} alkylNR⁵R⁶, where R^5 and R^6 are each independently H, C_{1-4} alkyl, aryl, or hetaryl, or may be joined to form an optionally substituted 3-8 membered ring optionally containing one of O, S or NR⁷;

wherein R^7 is H or C_{1-4} alkyl;

Q is CH;

W is C_{1-4} alkyl, or C_{2-6} alkenyl; where C_{1-4} alkyl or C_{2-6} alkenyl may be optionally substituted with C_{1-4} alkyl, OH, OC_{1-4} alkyl or NR¹⁵R¹⁶;

R^{15} , and R^{16} are each independently H or C_{1-4} alkyl, or may be joined to form an optionally substituted 3-8 membered ring optionally containing one of O, S or NR¹⁷;

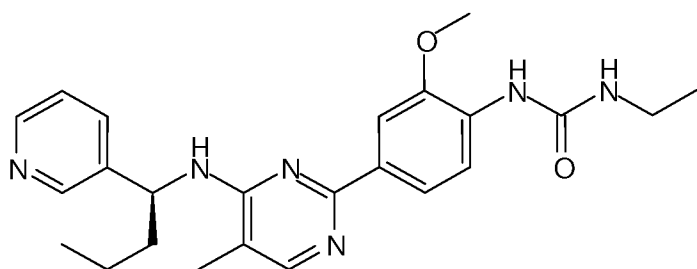
A is aryl, or hetaryl optionally substituted with 0-2 substituents independently chosen from halogen, C₁₋₄ alkyl, CF₃, aryl, hetaryl, OCF₃, OC₁₋₄ alkyl, OC₂₋₅ alkylNR¹⁸R¹⁹, Oaryl, Ohetaryl, CO₂R¹⁸, CONR¹⁸R¹⁹, NR¹⁸R¹⁹, C₁₋₄ alkylNR¹⁸R¹⁹, NR²⁰C₁₋₄ alkylNR¹⁸R¹⁹, NR¹⁸COR¹⁹, NR²⁰CONR¹⁸R¹⁹, and NR¹⁸SO₂R¹⁹;

wherein R¹⁸ and R¹⁹ are as defined in claim 10;

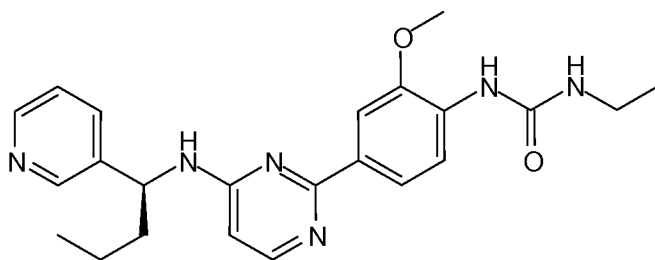
Y is selected from H, C₁₋₄ alkyl and NR²²R²³,

wherein R²² and R²³ are as defined in claim 10.

25. (new): The compound of claim 23 selected from:



and



or a pharmaceutically acceptable prodrug, salt, hydrate, solvate, crystal form or enantiomer thereof.

26. (new): A composition comprising a carrier and at least one compound according to claim 23.

27. (new): A composition comprising a carrier and at least one compound according to claim 24.

28. (new): A composition comprising a carrier and at least one compound according to claim 25.